PATENT COOPERATION TREATY

PCT

INTERNATIONAL SEARCH REPORT

(PCT Article 18 and Rules 43 and 44)

PCT/US05/35951		05 October 2005 (05.10.2005)	05 October 2004 (05.10.2004)
Applicant TOMOPHASE	CORPORATION		
This internation Basis of the a. With c c With reserved.	Article 18. A copy is being to that search report consists of the language, the important of the international attranslation of the of a translation further international search report authorized by or notified to the With regard to any nucleotide Certain claims were found to the title, the text is approved as submitted.	nternational search was carried out on the basis application in the language in which it was file international application into	is of: d, which is the language h (Rules 12.3(a) and 23.1(b)) e rectification of an obvious mistake
to the second of	ard to the drawings , gure of the drawings to be properties as suggested by the as selected by this A	according to Rule 38.2(b), by this Authority a the date of mailing of this international search ublished with the abstract is Figure No. 26	report, submit comments to this Authority

Form PCT/ISA/210 (first sheet) (April 2007)

INTERNATIONAL SEARCH REPORT

International application No.

PCT/US05/35951

A. CLASSIFICATION OF SUBJECT MATTER IPC: A61B 5/00(2006.01),6/00(2006.01)				
USPC: 600/310,316,322,365,473,476 According to International Patent Classification (IPC) or to both national classification and IPC				
B. FIELDS SEARCHED	······································	· · · · · · · · · · · · · · · · · · ·		
Minimum documentation searched (classification system followed U.S.: 600/310,316,322,365,473,476	by classification symbols)			
Documentation searched other than minimum documentation to the	e extent that such documents are included in	the fields searched		
Electronic data base consulted during the international search (nam	e of data base and, where practicable, search	n terms used)		
C. DOCUMENTS CONSIDERED TO BE RELEVANT				
Category * Citation of document, with indication, where a	appropriate, of the relevant passages	Relevant to claim No.		
X US 6,725,073 B1 (MOTAMEDI et al) 20 April 2004 column 3, lines 3-12 and 34-36, column 6, lines 3-1 1-7, figure 10 and claims 4, 38 and 44	· ·	10-17		
	US 6,725,073 B1 (MOTAMEDI et al) 20 April 2004 (20.04.2004), abstract, column 1, lines 1-9			
	6-14, columns 2, lines 49-57, column 3, lines 5-10, column 7, lines 57-64 and figure 10 US 5,803,909 (MAKI et al) 8 September 1998 (08.09.1998), column 1, lines 6-14, column 1-9, 18-20 16, lines 8-15			
Further documents are listed in the continuation of Box C.	See patent family annex.			
* Special categories of cited documents: "A" document defining the general state of the art which is not considered to be of particular relevance	"T" later document published after the inter date and not in conflict with the applica principle or theory underlying the inven	tion but cited to understand the		
"E" earlier application or patent published on or after the international filing date	"X" document of particular relevance; the clean considered novel or cannot be considered when the document is taken alone			
"L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)	"Y" document of particular relevance; the cleonsidered to involve an inventive step combined with one or more other such	when the document is		
"O" document referring to an oral disclosure, use, exhibition or other means	being obvious to a person skilled in the			
"P" document published prior to the international filing date but later than the "&" document member of the same patent family priority date claimed				
Date of the actual completion of the international search	Date of mailing of the international search 2 9 AUG 2008	report		
12 June 2008 (12.06.2008) Name and mailing address of the ISA/US	Adthorized officer			
Mail Stop PCT, Attn: ISA/US Commissioner for Patents P.O. Box 1450	Brian Casler Telephone No. 5712723700			
Alexandria, Virginia 22313-1450 Facsimile No. (571) 273-3201				

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PATENT COOPERATION TREATY

From the INTERNATION	NAL SEARCI	HING AUTHO	ORITY			
To: BING AI 12390 EL CAMINO REAL		PCT				
SAN DIEGO, CA 91230		WRITTEN OPINION OF THE INTERNATIONAL SEARCHING AUTHORITY				
					(PCT Rule 43bis.1)	
				Date of mailing (day/month/year)	2 9 AUG 2008	
Applicant's or	r agent's file re	eference		FOR FURTHER	RACTION	
17370-006WC	DI				See paragraph 2 below	
International a	application No		International filing date	(day/month/year)	Priority date (day/month/year)	
PCT/US05/35	951		05 October 2005 (05.10	2005)	05 October 2004 (05.10.2004)	
International F	Patent Classifi	cation (IPC) o	r both national classificat	ion and IPC		
	B 5/00 (2006./310,316,322,3	•	6.01)			
Applicant						
TOMOPHASI	E CORPORA	TION		· · · · · · · · · · · · · · · · · · ·		
1. This opin	ion contains in	ndications rela	ting to the following item	s:		
В	ox No. I	Basis of the	opinion			
Box No. II Priority						
Вс	ox No. III	Non-establis	hment of opinion with re	gard to novelty, inve	entive step and industrial applicability	
Box No. IV Lack of unity of invention			y of invention			
Bc	Box No. V Reasoned statement under Rule 43bis.1(a)(i) with regard to novelty, inventive step or industrial applicability; citations and explanations supporting such statement			•		
Box No. VI Certain documents cited			iments cited			
Box No. VII Certain defects in the international app		oplication				
Вс	ox No. VIII	Certain obse	rvations on the internatio	nal application		
2. FURTH :	ER ACTIO	N				
Internation Authority	nal Prelimina other than th	ry Examining is one to be the	Authority ("IPEA") ex	cept that this does IPEA has notified t	be considered to be a written opinion of the sonot apply where the applicant chooses and the International Bureau under Rule 66.1 bis(b) lered.	
IPEA a w of Form P	ritten reply to CT/ISA/220 c	gether, where or before the ex	appropriate, with amend xpiration of 22 months from	ments, before the ex	PEA, the applicant is invited to submit to the spiration of 3 months from the date of mailing whichever expires later.	
For furthe	er options, see	Form PCT/IS/	A/220 .			
3. For furthe	r details, see n	notes to Form l	PCT/ISA/220.			
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Name and mai	iling address o Stop PCT, Attn:		Date of comple	tion of this opinion	Authorized officer	
Comn	nissioner for Pa		17 June 2008 (1	7.06.2008)	Brian Casler	
Alexa	P.O. Box 1450 Alexandria, Virginia 22313-1450 Facsimile No. (571) 273-3201			Telephone No. 5712723700		

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International application No.	
PCT/US05/35951	

Box No.	I Basis of this opinion
 	gard to the language, this opinion has been established on the basis of:
	the international application in the language in which it was filed
i i	translation of the international application into, which is the language of a translation furnished for the purposes of nternational search (Rules 12.3(a) and 23.1(b)).
	This opinion has been established taking into account the rectification of an obvious mistake authorized by or notified to this Authority under Rule 91 (Rule 43bis.1(a))
	egard to any nucleotide and/or amino acid sequence disclosed in the international application, this opinion has been hed on the basis of:
a. 1	ype of material
	a sequence listing
[table(s) related to the sequence listing
b. 1	format of material
	on paper
] [in electronic form
c. t	ime of filing/furnishing
[contained in the international application as filed.
	filed together with the international application in electronic form.
	furnished subsequently to this Authority for the purposes of search.
	n addition, in the case that more than one version or copy of a sequence listing and/or table(s) relating thereto has been filed or furnished, the required statements that the information in the subsequent or additional copies is identical to that in the application as filed or does not go beyond the application as filed, as appropriate, were furnished.
5. Addition	nal comments:
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Box No. V Reasoned statement under Rul applicability; citations and exp		-	ep or industrial
1. Statement			
Novelty (N)	Claims	1-9,11-14 and 18-20	YES
	Claims	10, 15-17	NO
Inventive step (IS)	Claims	NONE	YES
	Claims	1-20	NO
Industrial applicability (IA)	Claims	1-20	YES
	Claims	NONE	NO
2. Citations and explanations: Please See Continuation Sheet			

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INTERNATIONAL SEARCHING AUTHORITY
Supplemental Box In case the space in any of the preceding boxes is not sufficient.
V. 2. Citations and Explanations:
Claims 10 and 15-17 lack novelty under PCT Article 33(2) as being anticipated by US Patent No. 6,725,073 B1 to Motamedi (Motamedi).
In Reference to Claim 10
Motamedi teaches the use of an optical coherence tomography system for measuring analyte (i.e. glucose concentration). The system
consists of -
A device for optically measuring a sample (see figure 10), comprising:
a) a plurality of light sources emitting light at different wavelength bands centered at different wavelengths (see column 7, lines 57-64 and claim 4);
b) a single waveguide to receive and guide the light at the different wavelength bands in a first propagation mode (see 10 and claim 44);
c) a probe head coupled to the waveguide to receive the light from the waveguide and to reflect a first portion of the light back to the waveguide in the first propagation mode and direct a second portion of the light to a sample (see column 2, lines 49-59), the probe head

collecting reflection of the second portion from the sample and exporting to the waveguide the reflection as a reflected second portion in

d) an optical differential delay unit to produce and control a relative delay between the reflected first portion and the reflected second

e) a detection module to receive the reflected first portion and the reflected second portion in the waveguide and to extract information of

a second propagation mode different from the first propagation mode (see claim 38 and column 3, lines 3-12):

portion received from the single waveguide in response to a control signal (see figure 10 and column 6, lines 3-11);

the sample carried by the reflected second portion (see figure 10); and

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Supplemental Box

In case the space in any of the preceding boxes is not sufficient.

f) a control unit, which produces the control signal to the optical differential delay unit, to set the relative delay at two different bias values to select a layer of material inside the sample to measure an optical absorption of the selected layer at each and every wavelength from the different light sources (see column 9, lines 1-7).

In Reference to Claims 15-17

Motamedi has been shown to teach all of the limitations of claim 10. In addition Motamedi further teaches:

Re claim 15:

The device as in claim 10, further comprising an optical element to receive the light at different wavelength bands from the light sources and to combine the received light into the single waveguide. (see figure 10, and column 7, lines 57-64).

Re claim 16:

The device as in claim 10, wherein the light sources are fixed in their emitting wavelengths(see column 7, lines 54-56).

Re claim 17:

The device as in claim 10, further comprising a mechanism to move a position of the second portion of the light relative to the sample to measure different locations on the sample (see column 3, lines 6-8 and 34-36).

Claims 1-9, 11-13 and 18-20 lack an inventive step under PCT Article 33(3) as being obvious over US Patent No. 6,725,073 B1 to Motamedi (Motamedi) in view of US Patent No. 5,803,909 to Maki (Maki).

In Reference to Claims 1

Motamedi teaches:

A method, comprising:

- a) combining and guiding optical radiation from a plurality of light sources, each emitting at wavelengths within a spectral band different from others, towards a sample through a common optical waveguide (see figure 10, abstract, claims and 4 and column 7, lines 57-64);
- b) reflecting a first portion of the combined radiation away from the sample at its vicinity while directing a second portion of the combined radiation to reach the sample (see figures 10, and column 3, lines 5-9);
- c) collecting and guiding at least part of the reflected first portion and at least part of a reflected second portion from the sample towards a detection module through the common optical waveguide (see figures 10, and columns 2, lines 49-57 and 3, lines 9-10);

However, Motamedi does not explicitly disclose a method comprising:

- d) separating the light into a plurality of spectral bands corresponding to emitting spectral bands of the light sources; and directing light radiation of the separated spectral bands to a plurality of light detectors, respectively.
- Maki, in the same field of endeavor, discloses a multispectral optical system for imaging and measuring various metabolic parameters within the body (see figure 2 and abstract). Maki cites the implementation of a plurality of multispectral light irradiation and receiving units in order to measure the "interior of a living body with ease without adversely affecting the living body' (see column 1, lines 6-14).

Therefore, it would have been obvious to one of ordinary skill in the art at the time of the invention to have included the step of "separating the light into a plurality of spectral bands corresponding to emitting spectral bands of the light sources; and directing light radiation of the separated spectral bands to a plurality of light detectors, respectively" of Maki in the method of Motamedi in order to facilitate the measurement and imaging of in vivo information using light as explicitly taught by Maki (see column 1. lines 9-11).

In Reference to Claim 2

Motamedi in view of Maki has been shown to teach all of the limitations of claim 1. Motamedi further teaches:

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Supplemental Box

In case the space in any of the preceding boxes is not sufficient.

"The method as in claim1, further comprising controlling the reflected first portion to propagate inside the common waveguide in a first propagation mode and the reflected second portion to propagate inside the common waveguide in a second propagation mode that is different from the first propagation mode (see figure 10 and claim7).

Therefore, Motamedi in view of Maki teaches all claim 2 limitations.

In Reference to Claim 3

Motamedi in view of Maki has been shown to teach all of the limitations of claim 2. Motamedi further teaches:

"The method as in claim 2, wherein the first propagation mode is a first polarization mode of the common waveguide and the second propagation mode is the second polarization mode that is perpendicular to the first polarization mode of the common waveguide (see figure 10 and claim?).

Therefore, Motamedi in view of Maki teaches all claim 3 limitations.

In Reference to Claim 4

Motamedi in view of Maki has been shown to teach all of the limitations of claim 1. Motamedi in view of Maki further teaches:

"The method as in claim 1, further comprising using broadband light emitters as the light sources, wherein each emitter has a broad spectrum centered at a wavelength different from wavelengths of other emitters" (see Maki column 4, lines 7-14 and column 8, lines 1-10).

Therefore, Motamedi in view of Maki teaches all claim 4 limitations.

In Reference to Claim 5

Motamedi in view of Maki has been shown to teach all of the limitations of claim 1. Motamedi in view of Maki further teaches:

"The method as in claim 1, further comprising using tunable laser sources as the light sources, wherein each tunable laser source emits coherent light at a wavelength tunable through a spectral band that is different from spectral bands of other tunable laser sources" (see Maki column 8, lines 8-15).

Therefore, Motamedi in view of Maki teaches all claim 5 limitations.

In Reference to Claim 6

Motamedi in view of Maki has been shown to teach all of the limitations of claim 1. Motamedi in view of Maki further teaches:

"The method as in claim 1, further comprising: producing a delay between the reflected first and second portions; and modulating the relative delay between the reflected first portion and the reflected second portion to measure a variation in power of light at each light detector" (see Motamedi column 6, lines 5-11 and claim 44).

Therefore. Motamedi in view of Maki teaches all claim 6 limitations.

In Reference to Claim 7

Motamedi in view of Maki has been shown to teach all of the limitations of claim 1. Motamedi in view of Maki further teaches:

"The method as in claim1, further comprising: adjusting a relative delay between the reflected first portion and the reflected second portion received from the single waveguide at two different delay values to select a layer of material inside the sample to measure an optical absorption of the selected layer at each and every wavelength from the different light sources" (see Motamedi claim 44).

Therefore, Motamedi in view of Maki teaches all claim 7 limitations.

In Reference to Claim 8

Motamedi in view of Maki has been shown to teach all of the limitations of claim 1. Motamedi in view of Maki further teaches:

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"The method as in claim 1, further comprising simultaneously directing light from the different light sources through the single, common waveguide to the sample" (see Motamedi figure 10 and column 7, lines 57-64).

Therefore, Motamedi in view of Maki teaches all claim 8 limitations.

In Reference to Claim 9

Motamedi in view of Maki has been shown to teach all of the limitations of claim 1. Motamedi in view of Maki further teaches:

"The method as in claim 1, further comprising sequentially directing light from the different light sources, one at a time, through the single, common waveguide to the sample" (see Maki columns 4, lines 7-16 and 16, lines 3-15).

Therefore, Motamedi in view of Maki teaches all claim 9 limitations.

In Reference to Claim 11

receive and detector the different beams".

Motamedi has been shown to teach all of the limitations of claim 10. However, Motamedi fails to explicitly disclose:

"The device as in claim 10, wherein the detection module comprises: an optical device to separate light at different wavelength bands into different beams; and a plurality of optical detectors to respectively

Maki's implementation of a multispectral in vivo optical sensing imaging and measurement device with multiple light sources and detectors has already been discussed (see rejection for claim 1, limitation d; see also Maki figure 2 and column 8, lines 1-30). Once again, Maki asserts that the novelty of his invention is that it enables measurement of the "interior of a living body with ease without

Therefore, it would have been obvious to one of ordinary skill in the art at the time of the invention to have included the step "wherein the detection module comprises:

an optical device to separate light at different wavelength bands into different beams; and a plurality of optical detectors to respectively receive and detector the different beams" of Maki in the method of Motamedi in order to facilitate the measurement and imaging of in vivo information using light as explicitly taught by Maki (see column 1, lines 9-11).

In Reference to Claim 12

Motamedi has been shown to teach all of the limitations of claim 10. Motamedi further teaches:

The device as in claim 10, wherein the detection module comprises:

adversely affecting the living body' (see column 1, lines 6-14).

a) an optical device to convert a part of received light in the first propagation mode and a part of received light in the second propagation mode into light in a third

propagation mode that propagates along a first optical path and to convert remaining portions of the received light in the first and the second propagation modes into light in a fourth propagation mode that propagates along a second, different optical path (see figure 10, abstract and columns 2, lines 55-67);

b) a first optical element in the first optical path to separate light at different wavelength bands into a first set of different beams (see figure 10 and column 3, lines 3-11);

d) a second optical element in the second optical path to separate light at different wavelength bands into a second set of different beams(see figure 10 and column 3, lines 3-11);

However, Motamedi fails to explicitly teach the claim 10 device limitations below, but Maki does:

- c) a plurality of first light detectors to respectively receive and detect the first set of different beams from the first optical element (see figure 2, reference marks 8a-8c, and column 8, lines 1-30);
- e) a plurality of second light detectors to respectively receive and detector the second set of different beams from the second optical element (see figure 2, reference marks 8d-8f, and column 8, lines 1-30);

Maki cites reducing the measurement time and thereby lessening the test burden on a patient and improving overall system operating efficiency (see column 2, lines 44-48) as key benefits to this approach.

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Supplemental Box

In case the space in any of the preceding boxes is not sufficient.

Therefore, it would have been obvious to one of ordinary skill in the art at the time of the invention to have included the step "wherein the detection module comprises:

a plurality of first and second light detectors" of Maki in the method of Motamedi in order to facilitate the time to obtain a measurement and improve system efficiency as explicitly taught by Maki.

In Reference to Claim 13

Motamedi has been shown to teach all of the limitations of claim 11. Motamedi further teaches using an interferometer to effect the necessary signal conditioning(interference) needed between reference and measurement signals in order to image and measure the sample of interest. It is well known in the art that gratings offer an alternative to accomplish similar ends with regard to signal interference. Therefore, barring unexpected results, gratings would not offer any patentable distinction over interferometers and would be solely used as a matter of design choice.

Therefore, Motamedi in view of Maki teaches all claim 13 limitations.

In Reference to Claims 18-20

Claims 18-20 and corresponding device claims 10-12 differ only in the specification of the type of light source. In the case of claims 18-20, the light source is a plurality of tunable laser sources.

Maki further discloses that the light source may be from a "plurality of laser sources" as well (see column 16, lines 8-15).

Claim 14 lacks an inventive step under PCT Article 33(3) as being obvious over US Patent No. 6,725,073 B1 to Motamedi (Motamedi) in view of US Patent No. 6,377,840 B1 to Gritsenko (Gritsenko).